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10 - 16 - 02

GP 1646

THE UNITED STATES PATENT AND TRADEMARK OFFICE
(Case No. 98-385-J)

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PATENT

In re Application of: Hauptmann et al.)

Serial No.: 09/899,429)

Filed: July 3, 2001)

For: TNF Receptors, TNF Binding)
Proteins and DNAs Coding)
For Them)

Before the Examiner: E. O'Hara

Group Art Unit: 1646

Commissioner for Patents
Washington, D.C. 20231

Sir:

TRANSMITTAL LETTER

1. We are transmitting herewith the attached papers for the above-described patent application:
Response to Office Action and return postcard.
2. GENERAL AUTHORIZATION TO CHARGE OR CREDIT FEES: Please charge any additional fees or credit any overpayment to Deposit Account No. 13-2490.
3. CERTIFICATE OF MAILING BY "EXPRESS MAIL" UNDER 37 C.F.R. 1.10: The undersigned hereby certifies that this Transmittal Letter and the papers, as described in paragraph 1 hereinabove, are being deposited with the United States Postal Service with sufficient postage as "Express Mail Post Office to Addressee" in an envelope addressed to: Commissioner for Patents, Washington D.C. 20231, on October 15, 2002.

Respectfully submitted,

McDonnell Boehnen Hulbert & Berghoff

Dated: October 15, 2002

By: 

Donald Zuhn, Ph.D.

Reg. No. 48,710



IN THE UNITED STATES PATENT AND TRADEMARK OFFICE
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#13
10/28/02

Commissioner for Patents
Washington, D.C. 20231

Sir:

RESPONSE TO RESTRICTION REQUIREMENT MAILED SEPTEMBER 12, 2002

Responsive to the Restriction Requirement, mailed September 12, 2002, Applicants elect to prosecute those claims directed to methods for ameliorating the harmful effects of TNF in an animal, comprising administering to an animal in need of such treatment a TNF binding polypeptide of SEQ ID NO: 4 or SEQ ID NO: 6, designated as Group I by the Examiner, with traverse. The basis for Applicants' traversal of the requirement is as follows.

Applicants respectfully submit that there will be no undue hardship on the Office in performing a search with respect to methods for ameliorating the harmful effects of TNF in an animal, comprising administering to an animal in need of such treatment a TNF binding polypeptide of SEQ ID NOs: 2, 4, 6, 8, 12, 14, 16, 18, or 20. A ClustalW multiple sequence alignment of these polypeptides is shown in Exhibit A. The sequence alignment was performed using the application MacVector 7.1.1 (Accelrys, Cambridge, UK; <http://www.accelrys.com>) at the default settings. This sequence alignment indicates that there is a substantial degree of homology between the amino acid sequences set forth in SEQ ID NOs: 2, 4, 6, 8, 10, 12, 14, 16, 18, and 20.

The amino acid sequence of the TNF receptor protein is set forth in SEQ ID NO: 2 (specification p. 5, ln. 7-39). The amino acid sequence consisting of residues 41 to 201 of SEQ ID

NO: 2 (which is equivalent to the amino acid sequence set forth in SEQ ID NO: 4) encodes a secretable TNF-binding protein (specification p. 4, ln. 27-41). As shown in Exhibit A, the polypeptides set forth in 2, 4, 6, 8, 10, 12, 14, 16, 18, and 20 all possess this portion of the TNF receptor protein. Moreover, this portion constitutes between 76.3% (SEQ ID NO: 8) and 99.4% (SEQ ID NO: 6) of the polypeptides set forth in SEQ ID NOs: 2, 4, 6, 8, 10, 12, 14, 16, 18, and 20. With the exception of an addition of methionine residue at the 5' end of the polypeptides set forth in SEQ ID NOs: 6, 10, 16, and 20, the polypeptides set forth in SEQ ID NOs: 2, 4, 6, 8, 10, 12, 14, 16, 18, and 20 differ only by the presence or absence of sequences encoding the signal peptide (amino acid residues 1 to 29 of SEQ ID NO: 2; specification p. 21, ln. 35 to p. 22, ln. 1), the portion of pro-protein cleaved following secretion (amino acid residues 30 to 40 of SEQ ID NO: 2; specification p. 22, ln. 7-11), and the linker region (amino acid residues 202 to 211 of SEQ ID NO: 2; specification p. 22, ln. 12-15) of the TNF receptor protein. Applicants respectfully submit that there will be no undue hardship on the Office in performing a search with respect to methods for ameliorating the harmful effects of TNF in an animal, comprising administering to an animal in need of such treatment a TNF binding polypeptide of SEQ ID NOs: 2, 4, 6, 8, 12, 14, 16, 18, or 20, since a search for methods for ameliorating the harmful effects of TNF in an animal, comprising administering to an animal in need of such treatment a TNF binding polypeptide of SEQ ID NO: 4 or SEQ ID NO: 6 will identify all of the non-elected sequences.

Applicants do not believe any additional fee is required. However, the Commissioner is authorized to charge any deficiency to Deposit Account No. 13-2490. If Examiner O'Hara believes it to be helpful, she is invited to contact the undersigned representative by telephone at (312) 913-0001.

Dated: October 15, 2002

Respectfully submitted,
McDonnell Boehnen Hulbert & Berghoff

By: 

Donald Zuhn, Ph.D.
Reg. No. 48,710



EXHIBIT A

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ClustalW (v1.4) multiple sequence alignment

10 Sequences Aligned

Alignment Score = 54899

Gaps Inserted = 2

Conserved Identities = 161

Pairwise Alignment Mode: Slow

Pairwise Alignment Parameters:

Open Gap Penalty = 10.0 Extend Gap Penalty = 0.1

Similarity Matrix: blosum

Multiple Alignment Parameters:

Open Gap Penalty = 10.0 Extend Gap Penalty = 0.0

Delay Divergent = 40% Gap Distance = 8

Similarity Matrix: blosum

Processing time: 3.5 seconds

```
SEQ 2      1 MGLSTVPDLLLLPLVLLELLVGIYPSGVIGLVPHLGDREKRDSVCPQGKYI  50
SEQ 4      1                                     DSVCPQGKYI  10
SEQ 6      1                                     MDSVCPQGKYI  11
SEQ 8      1 MGLSTVPDLLLLPLVLLELLVGIYPSGVIGLVPHLGDREKRDSVCPQGKYI  50
SEQ 10     1                                     MLVPHLGDREKRDSVCPQGKYI  22
SEQ 12     1 MGLSTVPDLLLLPLVLLELLVGIYPSGVIG-----DSVCPQGKYI  39
SEQ 14     1 MGLSTVPDLLLLPLVLLELLVGIYPSGVIGLVPHLGDREKRDSVCPQGKYI  50
SEQ 16     1                                     MLVPHLGDREKRDSVCPQGKYI  22
SEQ 18     1 MGLSTVPDLLLLPLVLLELLVGIYPSGVIG-----DSVCPQGKYI  39
SEQ 20     1                                     MDSVCPQGKYI  11
                        *****
```

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SEQ 2      51 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL 100
SEQ 4      11 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL  60
SEQ 6      12 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL  61
SEQ 8      51 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL 100
SEQ 10     23 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL  72
SEQ 12     40 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL  89
SEQ 14     51 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL 100
SEQ 16     23 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL  72
SEQ 18     40 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL  89
SEQ 20     12 HPQNNSICCTKCHKGTYLYNDCPGPGQDTCRECESGSFTASENHLRHCL  61
                        *****
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SEQ 2      101 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 150
SEQ 4      61 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 110
SEQ 6      62 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 111
SEQ 8      101 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 150
SEQ 10     73 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 122
SEQ 12     90 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 139
SEQ 14     101 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 150
SEQ 16     73 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 122
SEQ 18     90 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 139
SEQ 20     62 SCSKCRKEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSENLFQCFNCSLCL 111
                        *****
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|--------|-----|--|-----|
| SEQ 2 | 151 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 200 |
| SEQ 4 | 111 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 160 |
| SEQ 6 | 112 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 161 |
| SEQ 8 | 151 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 200 |
| SEQ 10 | 123 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 172 |
| SEQ 12 | 140 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 189 |
| SEQ 14 | 151 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 200 |
| SEQ 16 | 123 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 172 |
| SEQ 18 | 140 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 189 |
| SEQ 20 | 112 | NGTVHLSCQEKQNTVCTCHAGFFLRENECVSCSNCKKSLECTKLCLPQIE | 161 |
| ***** | | | |

| | | | |
|--------|-----|---|-----|
| SEQ 2 | 201 | NVKGTEDSGTTVLLPLVIFFLGLCLLSLLFIGLMYRYQRWKSCLYSIVCGK | 250 |
| SEQ 4 | 161 | N | 161 |
| SEQ 6 | 162 | N | 162 |
| SEQ 8 | 201 | NVKGTEDSGTT | 211 |
| SEQ 10 | 173 | NVKGTEDSGTT | 183 |
| SEQ 12 | 190 | NVKGTEDSGTT | 200 |
| SEQ 14 | 201 | N | 201 |
| SEQ 16 | 173 | N | 173 |
| SEQ 18 | 190 | N | 190 |
| SEQ 20 | 162 | NVKGTEDSGTT | 172 |
| * | | | |

| | | | |
|--------|-----|--|-----|
| SEQ 2 | 251 | STPEKEGELEGTTTKPLAPNPSFSPTPGFTPTLGFSPVPSSTFTSSSTYT | 300 |
| SEQ 4 | 162 | | 161 |
| SEQ 6 | 163 | | 162 |
| SEQ 8 | 212 | | 211 |
| SEQ 10 | 184 | | 183 |
| SEQ 12 | 201 | | 200 |
| SEQ 14 | 202 | | 201 |
| SEQ 16 | 174 | | 173 |
| SEQ 18 | 191 | | 190 |
| SEQ 20 | 173 | | 172 |

| | | | |
|--------|-----|---|-----|
| SEQ 2 | 301 | PGDCPNFAAPRREVAPPYQGADPILATALASDPIPNNPLQKWEDSAHKPQS | 350 |
| SEQ 4 | 162 | | 161 |
| SEQ 6 | 163 | | 162 |
| SEQ 8 | 212 | | 211 |
| SEQ 10 | 184 | | 183 |
| SEQ 12 | 201 | | 200 |
| SEQ 14 | 202 | | 201 |
| SEQ 16 | 174 | | 173 |
| SEQ 18 | 191 | | 190 |
| SEQ 20 | 173 | | 172 |

| | | | |
|--------|-----|---|-----|
| SEQ 2 | 351 | LDTDDPATLYAVVENVPPLRWKEFVRRRLGLSDHEIDRLELQNGRCLREAQ | 400 |
| SEQ 4 | 162 | | 161 |
| SEQ 6 | 163 | | 162 |
| SEQ 8 | 212 | | 211 |
| SEQ 10 | 184 | | 183 |
| SEQ 12 | 201 | | 200 |
| SEQ 14 | 202 | | 201 |
| SEQ 16 | 174 | | 173 |
| SEQ 18 | 191 | | 190 |
| SEQ 20 | 173 | | 172 |

| | | | |
|--------|-----|--|-----|
| SEQ 2 | 401 | YSMLATWRRRTTPREATLELLGRVLRDMDLLGCLEDIEEALCGPAALPPA | 450 |
| SEQ 4 | 162 | | 161 |
| SEQ 6 | 163 | | 162 |
| SEQ 8 | 212 | | 211 |
| SEQ 10 | 184 | | 183 |
| SEQ 12 | 201 | | 200 |
| SEQ 14 | 202 | | 201 |
| SEQ 16 | 174 | | 173 |
| SEQ 18 | 191 | | 190 |
| SEQ 20 | 173 | | 172 |

| | | | |
|--------|-----|-------|-----|
| SEQ 2 | 451 | PSLLR | 455 |
| SEQ 4 | 162 | | 161 |
| SEQ 6 | 163 | | 162 |
| SEQ 8 | 212 | | 211 |
| SEQ 10 | 184 | | 183 |
| SEQ 12 | 201 | | 200 |
| SEQ 14 | 202 | | 201 |
| SEQ 16 | 174 | | 173 |
| SEQ 18 | 191 | | 190 |
| SEQ 20 | 173 | | 172 |